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Alkaloids of *Ptilularia sewerzowii*. S. Yuzusay, R. A. Konovalova and A. P. Orlikov. *J. Gen. Chem. (U. S. S. R.)*, 9, 1911-14 (1929).—Bruit with C_6H_5Cl of 100 g. of dry, powd. tubers of *Ptilularia sewerzowii* (Kavalenkov, 1900), family Liliaceae (gathered in August in Central Asia) gave 925 g. (0.92%) of crude bases which after complex process of reworking yielded a new alkaloid, named *alpinine*, $C_{21}H_{21}NO$, m. 271-2 $^{\circ}$, [α] D^{25} 108.5 $^{\circ}$; *HCl salt*, m. 332-3 $^{\circ}$; *Mel dextr.*, m. 310-11 $^{\circ}$. It contains 3 OH groups with a tertiary N and probably has the formula $C_{21}H_{21}(N)(OH)_3$. The investigation is being continued.
Chas. Blanc

430-314 METALLURGICAL LITERATURE CLASSIFICATION

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1. KONOVALOVA, P. A., YURIEV, S., OREKHOV, A.P.

2. USSR (600)

"On the Alkaloids of Plants of the Family Papaveraceae. Vi. The Alkaloids of the
Glacium Gibrilligerum", Zhur. Obshch. Khim., 9, No. 21, 1939. Alkaloid Dept.
All-Union Soci.-Res. Chomico-Pharmaceutical Inst. Imeni S. Ordzonikidze.
Received 4 June 1952.

9. [redacted] Report U- 1626, 11 Jan 1952

CA

Alkaloids of the family Papaveraceae. VII. Alkaloids of *Papaver armeniacum*. The structure of armepavine. S. Yumakov, R. A. Konovalova and A. P. Orlova. *J. Russ. Chem. (U. S. S. R.)* 10, 611-81(1940); cf. *C. A.* 34, 4072^{1,2}.—Armepravine (I), previously isolated from *Papaver armeniacum* and *Papaver floribundum* (*C. A.* 30, 1380³), yields on methylation with CH_3N_3 in MeOH and ether methyl-I (II), m. 63-4°, or -84.45° (in CHCl_3), the methiodide (III) of which m. 133 4°. II on boiling with Ag_2O followed by energetic oxidation with HNO_3 (d. 1.4) gives anisic acid. Heating of III with KOH and MeOH gives inactive *des-O,N-dimethylarmepavine* (IV), m. 80-7°; IV- HCl , m. 229-30°; IV-MeI (V), m. 233-4°. On heating V with KOH and MeOH , MeI is split off with the formation of a compd. $\text{C}_{10}\text{H}_{16}\text{O}_2$, (VI), m. 79-9.5°. Oxidation of VI by means of KMnO_4 in acetone yields anisic acid and 4,5-dimethoxyphthalic acid. This shows that VI is 4,5,4'-trimethoxy-2-vinylstilbene and I is 6,7-dimethoxy-1-(4-methoxybenzyl)-2-methyl-1,2,3,4-tetrahydroisoquinoline. Ethylation of I with Et_2SO_4 and NaOH gives *diethyl-des-armepavine* [*dimethoxyphenyl-2-(β-ethylmethylaminomethyl)stilbene*] as a thick oil, which on

oxidation with KMnO_4 in acetone yields *p*-ethoxybenzoic acid. This result proves that I is 6,7-dimethoxy-1-(4-hydroxybenzyl)-1-methyl-1,2,3,4-tetrahydroisoquinoline, which conclusion is further substantiated by the oxidation of I with KMnO_4 in acetone whereby 6,7-dimethoxy-1- *keto*-2-methyl-1,2,3,4-tetrahydroisoquinoline, m. 124-5°, is obtained.

Gertrude Berend

ASA-11A - METALLURGICAL LITERATURE CLASSIFICATION

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| 153 | 3 | | | | | | |

YUNUSOV, S.

Yunusov, S. - "On the dynamics of accumulation, the role and the formation of alkaloids in plants", Izvestiya Akad. nauk UzSSR, 1948, No. 4, p. 11-27, (Resume in Uzbek), Bibliog: 67 items.

SO: U-3042, 11 March 1950, (letopis 'nykh Statey, No. 10, 1949).

YUNUSOV, S. Yu.

Yunusov, S. Yu. "On determining the structure of alkaloids of the aporphine group," Doklady Akad nauk UzSSR, No. 8, 1948, p. 12-16 - Resume in Uzbek language - Bibliog; p. 16

SO: U-3850, 16 June 53, (Letopis 'Zhurnal 'nykh Statey, No. 5, 1949).

YUNUSOV, S.

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USSR/Chemistry - Alkaloids Mar 1948
Chemistry - Alkaloids, Zengoricum

"The Alkaloids Aconitum Scengoricum Stapf. I.
Alkaloids of Ranunculaceae Family," S. Yunusov, Lab
of Chem of Alkaloids, Inst Chem, Acad Sci Uzbek SSR,
Tashkent, 12½ pp

"Zhur Obshch Khim" Vol XVIII (LXXI), No 3

New alkaloid having formula $C_{21}H_{29}NO_3$, known as
'zengorinon,' was obtained from roots of Aconitum
scengoricum Stapf. Gives description of following
'zengorinons': semicarbazone, diastyl-, dihydro-,
dihydrodiacetyl-, dichlor-, iodomethylate-zengorines.
Submitted 28 Jan 1947.

69T6

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The alkaloids of *Lemnica ewersmannii* and *Leontice alberti*. S. Yunusov and L. G. Sorokina (Acad. Sci. U.S.S.R.), *J. Gen. Chem. (U.S.S.R.)* 19, No. 10, 3427-35 (1949) (English translation).--See *C.A.* 44, 1997b.
B. J. C.

CA (UNUSC, p.

Alkaloids of *Cocculus laurifolius*. I. S. Yunusov
(Lab. Alkaloid Chem., Inst. Chem., Acad. Sci. Uzbek.
S.S.R., Tashkent). *Zhur. Obshchey Khim.* (J. Gen.
Chem.) 20, 368-75 (1950).—Extr. of *Cocculus laurifolius*
leaves with $(CH_3)_2Cl$, in NH_4OH (5%) gave 0.9% total
alkaloids. The nonphenolic fraction treated with HI in
 Me_2CO gave, from 45 g. crude oil, 41 g. cocculidine-HI,
m. 174-5° (from Et₂OAr after drying at 90-100° in vacuo);
the free base liberated by NH_4OH , m. 28-7° (from petr.
ether), is $C_{18}H_{20}O_2N$, [α]_D 250.9° (in $CHCl_3$), having no
 N Me or OMe groups; *m.p.* m. 131.5-2.5° (crude oil),
m. 137-8° (from Me_2CO); *methiodide*, m. 238-9°. The
phenolic fraction of alkaloids crystd. from Me_2CO gave
(from 8 g. crude oil) 3.8 g. *recrystd.* $C_{18}H_{20}O_2N$, m. 217-
18° (from Me_2CO), *m.p.* 271.1° ($Mg(OH)_2$); *nitrile*, m.
196-7° (from EtOH); *HCl salt*, m. 222-3°; methylation
with CH_3N_3 gave the $-OMe$ deriv., identical with the
methiodide of cocculidine (mixed m.p.), i.e., dimethoxy
deriv. of $C_{18}H_{20}N$. G. M. Kosolapoff

Alkaloids of Ranunculaceae. III. Alkaloids of Thalictrosum mians. S. Yumusov and N. N. Progessor. (Lab. Khim. Alkaloid. Inst. Khim. Akad. Nauk Uzbek. S.S.R., Tashkent). Zher. Obshch. Khim. (J. Gen. Chem.) 20, 1151-61 (1950); cf. C.A. 44, 7025n. —Extr. of air-dried *Thalictrosum mians.* (with CH_3Cl) in the presence of 10% NaOH gave 0.31% total alkaloids. Extr. with EtOH of 121.2 g. of such a mixt. gave 40 g. insol. thalicmine, $\text{C}_{11}\text{H}_{15}\text{NO}_3$, decompr., 237-4° (crude), m. 233° (pure, from $\text{EtOH}-\text{CHCl}_3$), $[\alpha]_D^{25} -64.5^\circ$; HCl salt, decompr. 147-57° (from EtOH); methiodide, decompr., 250° (from EtOH); perchlorate, decompr., 238-41°. The alc.-sol. alkaloid portion yielded on cooling to -10° 14 g. thalidazine, $\text{C}_{11}\text{H}_{14}\text{N}_2$, decompr., 102-3° (from EtOH), $[\alpha]_D^{25} 232.2^\circ$ (CHCl_3); methiodide, decompr., 234-5°. Similar extr. of the roots of the plant gave 0.95 or 0.83% total alkaloids, depending on the time of collection (spring or fall). These yielded 3 alkaloids: thalicmine, $\text{C}_{11}\text{H}_{15}\text{NO}_3$, m. 137.8° (from MeOH), isolated through the poorly sol. HCl salt, m. 238-70°, $[\alpha]_D^{25} 255.3^\circ$ (EtOH); *III* salt, decompr., 229-4° (in sealed tube); HBr salt, m. 238-60° (from H_2O); methiodide, m. 239-7° (in sealed tube); acetylation of this alkaloid with AcCl 12 days at room temp. gave an *Ac* deriv. m. 191-2°, which is optically inactive. The mother liquor after isolation of thalicmine yields upon neutralization with NH_3 and treatment with tartaric acid (in EtOH) 17 g. thalidimine borate, decompr., 239-40° (in sealed tube); free base, $\text{C}_{11}\text{H}_{15}\text{NO}_3$, m. 192-3°, $[\alpha]_D^{25} -54^\circ$ (EtOH), with neg. reactions for aldehyde, ketone, or methylenedioxy groups; *III* salt, decompr., 222-6° (sealed tube); methiodide, m. 217-17.5° (from EtOH). Extr. of the residual alkaloid mass with hot EtOH and Me_2CO gave an orange base, m. 243°. All 3 alkaloids appear to be derivs. of 2-methylbenzyltetrahydroisoquinoline, the latter being the structure of thalicmine.

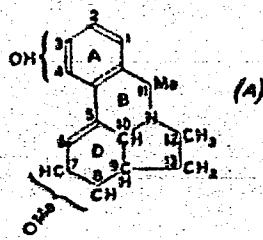
(S. 31. Konolopoff)

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Alkaloids of *Cocculus laurifolius*. II. Structure of coccidine and cocculine. S. Yunusov (Lab. Khim. Alkaloid. Inst. Khim. Akad. Nauk UZSSR, S.S.R., Tashkent), Zhur. Osnov. Khim. (I. Gen. Chem.) 20, 1514-23 (1950); cf. J. Am. Chem. Soc. 72, 6822. The 1st step of the Holmann degradation of cocculine with KOH and AgOH provides similar to that of 1-substituted tetrahydroisoquinoline alkaloids yielding optically active *dextro*-N-methylcoccidine, m. 82.3° (from petr. ether), [α]_D 108.80°; *enantiomeric*, m. 205.6° (from MeOH). The B ring is opened with loss of asymmetry at C atom 11. The 2nd step of the degradation gives a double bond largely at the 9,10-position (ring D), followed by loss of a MeO group as MeOH, with aromatization, leading to *dextro*-N-methylcoclaurine, C₁₇H₂₁NO, b. 160.8°, optically inactive, and conig. 1 MeO group and 1 vinyl link, *m.p.* 117, m. 195.0°. The 3rd step of the degradation yields *ortho*-*nitro*-*2,2'-dimethoxy*, b. 218°, which with KMnO₄ in NaClO gives *ortho*-*nitro*-*2,2'-diphenylbenzoic acid*, m. 180-90° (from Et₂O), optically inactive; *dextro*-ester, m. 73-4° (from petr. ether). Distill. of the acid with Zn dust gave *fluorescein*. Hence the N of cocculine and coccidine is at the fusion point of the pyridine and piperidine rings; the phenolic group in cocculine is in the aromatic ring A, while the double bond and MeO are in ring D, the possible location of the double bond being at 5,6 or 7,8. Cocculine and cocculidine are derivs. of phenanthridine and indole. The plant also contains a 3rd alkaloid of nonphenolic type, which is a deriv. of 1-methyl- or 1-benzyltetrahydroisoquinoline. G. M. Kosolapoff

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Alkaloids of *Cocculus laurifolius*, D.G. II. Structure of coccinaline and cocconine. N. Nunnally. (*J. Am. Chem. USSR*, 1939, 22, 1514-1523 [U.S. transl., 1977-1986]).—The first stage of the Holmann degradation of coccinaline, under the action of KOH or AgOH, takes place in the same manner as with 1-substituted tetrahydroisoquinoline alkaloids, with the formation of optically active "d-N-methyl-coccinaline". In this reaction, the chief result is the opening of ring B and the destruction of the asymmetry of C₁₀. At the second stage of the degradation, carried out by the action of KOH or AgOH, a double linking is formed chiefly between C₁₀ and C₁₂. After the appearance of this second double linking in ring D, MeOH is eliminated by union of OM₂ with H, and the ring acquires aromatic character. Thus, after the second stage of the degradation, all four C atoms have lost their asymmetry, and optically inactive "d,N-dimethylcoccinaline" is obtained containing 1 OM₂ and 1 CH₂CH₃ group. The third stage of the degradation leads to the formation of methoxy-2 : 3'-divinyl biphenyl, the structure of which is



Area

YUNUSOV, S.

"The alkaloids of Delphinium semibarbatum. IV. Alkaloids of the Ranunculaceae order." by S. Yunusov and N. K. Abubakirov. (p.174)

SO: Journal of General Chemistry (Zhurnal Obshchei Khimii) 1951, Volume 21,
No. 1.

CA

Alkaloids of the Ranunculaceae family. V. Delphinium
oreophilum. S. Yannayev and N. K. Abubakirov. *J.
Gra. Chem. U.S.S.R.* 21, 1059-63 (1931) (Engl. transla-
tion).—See C.A. 26, 5168.

Card 1 of 1

USSR Chemistry - Alkaloids

Jun

"Alkaloids of Thalictroides L. II. The Structure of Thalicmine and Thalimidine," S. I. Issov, N. N. Progessov, Lat. of Alkaloid Chem., Inst. of Chem., Acad. Sci. Uzbek SSR, Tashkent, Zhur Obsch Khim" Vol XXII, No 6, pp 1047-1051.

The formula of thalicmine $C_{20}H_{23}NO_4$ as $C_{16}H_{10}(NCH_3)(OCH_3)_3(OH)$. The hydroxyl group has weakly phenolic properties. The OH group retains these properties in the dea-N-methylthalicmine. Oxidation of the substance obtained by boiling

218T?7

USSR/Chemistry - Alkaloids (Contd 1) Jun 5,2

thalicmine in acetic anhydride yields melophanic acid. Oxidation of thalicmine itself yields meta-hemipinic acid. Hofmann's degradation of O-methyl-thalicmine, repeated twice, yielded a nitrogen-free substance, which upon oxidation yielded tetra-methoxy-phenanthrene-carboxylic acid. The roots of Thalictrum minus L., in the immature state, contain D-glucosine, which was identified as O-methyl-thalimidine. The structure of thalicmine is that of a 2,3,5-trimethoxy-6-hydroxyporphine. Melophanic acid and phenanthrene were obtained from N-acetylthalicmine, which is optically inactive and not basic. After Hofmann's degradation had

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(CRA 47 no.16: 8084 J3)

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USSR/Chemistry - Alkaloids (Contd 2)

been carried out twice, thalicmine yield trimethoxy methylene dioxy-vinylphenanthrene. Thalicmine is a deriv of pentahydroxyaporphine. Apparently its structure corresponds to 3,4,7-trimethoxy-5,6-methyl-enedioxy aporphine.

218T27

IR Chem - Alkaloids

Jun

of Acad. M. V. Lomonosov, Institute of Chemistry,
V. I. Vernadsky Institute of Geochemistry and
E. N. Saryakina, Institute of Chemistry,
Yunusov, Inst. of Chem., Acad. Sci. Uzbek SSR, Tashkent

but Ovchinnikov, Khim Vol XXII, No 6, pp 1054-1061

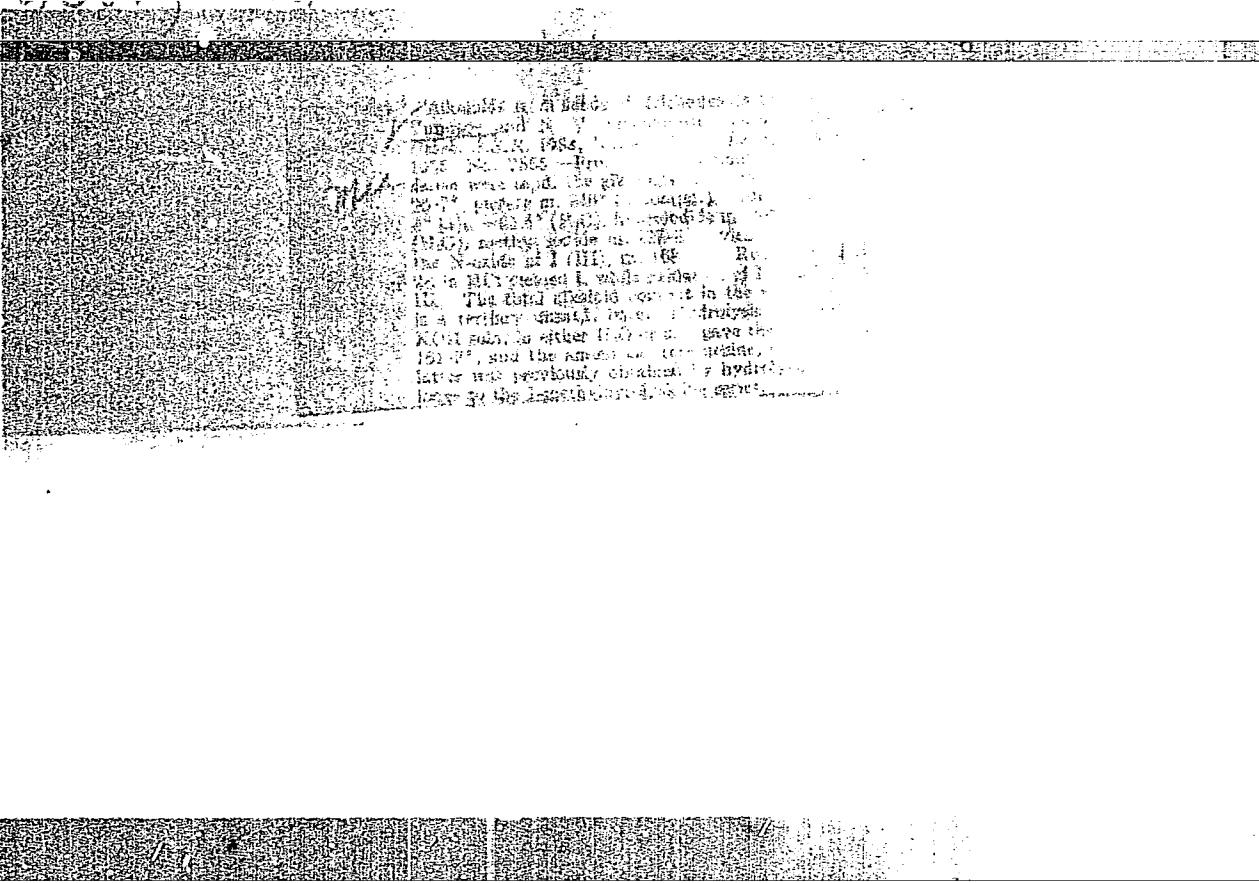
obtained the following alkaloids from the leaves,
stems, and young stems of Haplophyllum perforatum
(M.B.) Kar. et Kir.: ocimannanine, a cryst base with
mp 110-111°, and the new alkaloid haploperine
 $C_{17}H_{25}NO_4(OCH_3)_2$, mp 155-156°. Obtained the hydro-
chloride or haploperine (mp 129-131°) and hexahy-
drohaploperine $C_{17}H_{25}NO_6$ (mp 159-160°). Haploperine
218gr28

JSER/Chemistry - Alkaloids (Contd)

Jun 52

reacts with acids to form $C_{17}H_{17}NO_5$, mp 138-139°.
Under the action of CH_3I it forms the isomeric complex
 $C_{17}H_{25}O_4(=CO)(=N-CH_3)(OCH_3)$. It obviously is a
quinoline deriv. The alkaloids are contained mostly
in the leaves and seeds. Investigated the alkaloid
content of the above plants.

218gr28



U.S. No. 5
Mar. 30, 1954
Organic Chemistry

YUNUSOV, S.

The alkaloids of Thallium minus. II. The structure of
thallicine and thalimine. S. Yunusov and N. N. Pro-
gresov (Azer. Sci. Uzbek. Tashkent). J. Gen. Chem.
U.S.S.R. 22, 1095-1101 (1952) (Engl. translation).—See
C.A. 47, 8084i. (3)

AK
1-26-51

YUNUSOV, I.S.

Chemical Abst.
Vol. 48 No. 5
Mar. 10, 1954
Organic Chemistry

The alkaloids of Haplophyllum perforatum, H. pedicellatum, H. dubium, H. bucharicum, and H. versicolor. I. S. Yunusov and G. P. Sidiyakin (Acad. Sci. Uzbek, Tashkent). *J. Gen. Chem. U.S.S.R.* 22, 1103-8 (1952) (Engl. translation).—See *C.A.*, 47, 8084f.

H. L. H.

MF
1-26-54

YUNUSOV, S.

Chemical Abst.
Vol. 48 No. 5
Mar. 10, 1954
Organic Chemistry

Alkaloids of Delphinium. IV. Delsemidine. S. Yunusov and N. K. Abubakirov (Acad. Sci. Tashkent). J. Gen. Chem. U.S.S.R. 22, 1503-5 (1952) (Engl. translation).
See C.A. 47, 7515i.

H. L. H.

MF
1-285A

YUNUSOV, S.YU.

The Committee on Cultural Prizes for the Journal of Ministry of Finance of the Uzbek SSR scientific and technical press informed me that the following scientific works, popular scientific books, and textbooks have been submitted for competition for Stalin Prize for the year 1958 (Sovetskaya Kultura, Moscow, No. 28-29, 20 Feb - 2 April 1959).

| Name of author | Title of work | Organization |
|----------------|--|--|
| Yunusov, S.Yu | "Investigation in the Field of the Chemistry of Alkaloids" | Presidium, Academy of Sciences Uzbek SSR (6) |

22 APRIL 1959

YURKOV S. M. and ABDUBAKIROV N. K.

Delbine and Delphatine. V. Study of Delphinium Alkaloids, Page 1446,
Sbornik stately po obshchey khimii (Collection of Papers on General
Chemistry), Vol II, Moscow-Leningrad, 1953, pages 1680-1686.

Laboratory of the Chemistry of Alkaloids, Inst of Chemistry, Acad
Sci Uz SSR

U S S R .

✓ Delphinium *Adonis* -
V. K. Abrosimov
Prod. Tashkent, USSR
Sesquiterpenes
Lactone
Lactone of 1,4-diene-3,5-diol
It is followed by ketone then
ketone epoxide and lactone
acid. On this basis it can be assumed that the
product is

$\text{C}_{16}\text{H}_{20}\text{O}_4$. It is believed that this product
Delphinium contains 1,4-diene-3,5-diol which is
phloracetophenone derivative. The presence of
hydroxyl of delphinium is supported by
decolor 210°
100°, Melting point 160°
It is soluble in
solvent of benzene, chloroform, ether,
etc. KMnO₄ gives
Mel in MeOH
presence of
dihydroxyacetone
ketone
ketone epoxide
acid

"APPROVED FOR RELEASE: 03/15/2001

CIA-RDP86-00513R001963120020-7

APPROVED FOR RELEASE: 03/15/2001

CIA-RDP86-00513R001963120020-7"

"APPROVED FOR RELEASE: 03/15/2001

CIA-RDP86-00513R001963120020-7

~~100% ¹⁸O in a dilute solution of H₂O (III), m. 70-80°~~

~~H₂O (III) is a dilute solution of H₂O (III), m. 70-80°~~

APPROVED FOR RELEASE: 03/15/2001

CIA-RDP86-00513R001963120020-7"

11/11/507, S. G.

Investigation of alkaloids from *Aconitum heterophyllum*
S. Yu. Yunusov, E. V. Sretkova and G. I. Dzhankina
Doklady Akad. Nauk Uzb. SSR 1954, No. 21, p. 211
Referat. Zhur. Khim. 1954, No. 49(14), p. 1150 (1954).

The roots of this plant are used in central Asia as a medicament. The above-ground part contains 1.9% and the roots 1.02% of alkaloids. To isolate the alkaloids the aqueous ethanolic ext. was treated with dil. acid and the mixt. of bases in the acid ext. was precip. with NH_4OH . The mother liquor was satd. with NaCl and ext'd. with CHCl_3 . The ext. was fractionated with org. solvents grad. decreasing in the strength of the bases up to 20% of cryst. alkaloids were obtained. Acid or alk. vapor of the mixt. followed by fractionation yielded 50% of cryst. alkaloids and various salts. The largest compound was the optically inactive dihydro-*taizamine*, m. 145-5°. Its formula is $\text{C}_{11}\text{H}_{14}\text{N}_2$ and not as previously suggested, $\text{C}_{11}\text{H}_{14}\text{N}^+$ (cf. Konon'cov and Orekhov, *Zhur. Obshch. Khim.* 10, 745 (1940)). The tertiary N atom is combined with an Et radical. Oxidation with KMnO_4 liberated MeCHO . The alkaloid contains 3 OMe and 2 OH groups. Cryst. monoacetate and acetate were also obtained. A second alkaloid was apparently *taizamine*, $\text{C}_{11}\text{H}_{14}\text{N}_2$, m. 246-7°, $\alpha_{D}^{25} +35.6^\circ$. It contained a double bond and had no OMe groups. A third alkaloid, m. 241-2°, $\alpha_{D}^{25} -85.1^\circ$.

YUNUSOV, S. Yu.

USSR/Chemistry

Card 1/1

Authors : Abubakirov, N. K.; and Yunusov, S. Yu.
Title : Investigation of Delphinium alkaloids. Part I. Structure of delphisine.
Periodical : Zhur. Ob. Khim. 24, Ed. 4, 737 - 747, April, 1954.
Abstract : The author carried out the transformation of the delphisine nitroso compound into the nor-base. Treatment of the latter with iodide produces a compound identical with delphisine. The presence of the N-n-butyl group is decisive for the course of the conversions. Fifteen references; 10 Japanese, English since 1934.
Institution : Institute of Chemistry at the Academy of Sciences of the
Submitted : December 11, 1957.

Yanusov, S. Yu.

62 *V. Alkaloids of Aconitum taizhouense* S. Yu. Yanusov, V. P. Schikova, and G. F. Pudovkin *Zhur. Org. Khim.* 1974, 24, 2237-42 (1984), ref. 42, 7947. - 1) *n*-HCl ext. of the crude alkaloid mass obtained from 22.9 mg of dried plant leaves and stems was reduced 5 hrs with 10% H₂S₂O₄, cooled, ext'd with Et₂O, filtered, and treated with NaCl. The pfd material was ext'd with CH₂Cl₂ and the ext evapd and rubbed with MeOH giving 1.2 g of product (I), with further amts recovered from the ext; the total yield was 92% λ . The Et₂O ext yielded a very small amount (0.1%). Fractional pptn of the crude product of I was done from 10% H₂S₂O₄ by gradual addition of NaCl. This ext, after treatment with 1M NaOH, gave 10.9% I and a long-chain saponine, I, m. 145.0° (from MeOH); *n*-C₁₀H₂₀O₂; *H*P₂O₄, m. 78-80° (from BuOAc). II, salt-salt, m. 67.5-8.5°, $[\alpha]_D^{25} -12.8^{\circ}$ (H₂O); (anhyd. salt, dioxane, 120°, $[\alpha]_D^{25} -14.8^{\circ}$ (H₂O)). Some decomp. m. 93.5° (from standing with Ac₂O) and yields the d. salt, m. 117-9°, $[\alpha]_D^{25} -5.35^{\circ}$. MeOH-hydrolysis of this salt w/ NaOH gave the original alkaloid, I, MeOH, m. 223.1°. Oxidation of I with KMnO₄-H₂SO₄ gave Acid. Thus I can be represented by *C₁₈H₂₄NiKOH₂·MeOH*. Takizawa m. 116.7° (ref. 35, 5° (cf. Yanusova and Grigor'ev, *Zh. Org. Khim.* 1974, 10, 34-45)). *n*-C₁₀H₂₀O₂ has 1 double bond as shown by IR spectrum (KBr, m. 3000-3500 cm⁻¹); *n*-C₁₀H₂₀O₂ has 1 double bond as shown by IR spectrum (KBr, m. 3000-3500 cm⁻¹).

✓Alkaloids of seeds of Lolium canescens S. Yu. Yunusov
and S. F. Akramov *Zhur. Obshchel. Khim.* 23, 1818-20
(1953). - The $(\text{CH}_2\text{Cl})_2$ ext. of the seeds of *L. canescens*
gave a CHCl_3 -sol fraction of alkaloids which made up
0.26% yield on seed wt. The following alkaloids were
isolated by fractional distn.: *lepidine*, *stachydrine*, *loline*,
loline (10^4), which can be sublimed (HCl salt,
 $(\text{CO})(\text{NMe})(\text{NH})$, m. 210-12°); *loline* ($1.$), *loline*
(α) 0.2%, used for purification of the free base, b.p. 229°,
66.1-434.0, n_D²⁰ 1.6603, [α]_D²⁰ 18.0° [*d*-*H* salt], decamp.
254.5°, [α]_D²⁰ 3.6°, *d*-*H* salt, decamp. 253.5°, [α]_D²⁰ 0.4°;
loline m. 148.9°, [α]_D²⁰ 0.18°; *mononitrate*, m. 219.5°,
parahydronate, decamp. 282°; *sulfate*, decamp. 360.5°; *d*-
loline, decamp. 258.6°; *loline* m. 219.12° (from dil.
MeOH); is accompanied by *N-methyl-loline* *Mel*, decamp.
267.9°, [α]_D²⁰ -4.84° on treatment of 1 with *Mel*. The
latter is sepd. by virtue of greater solv. in dil. MeOH
[0.2 g.] with 0.98 g. 40% formalin and 0.99 g. HgCl_2Hg re-

fluxed 8 hrs. Freezing g. *N-methyl-loline* m. 219.12°

n_D²⁰ 1.6602, [α]_D²⁰ 0.13°

9.5% *HgCl* salt, decamp. 267.9°, [α]_D²⁰ 0.13°

[α]_D²⁰ 6.1°, decamp. m. 219.12°

Treatment of 1 with HgCl_2Hg and NaNO_2 gave with 0.98 g.

loline m. 219.12°, [α]_D²⁰ 0.13°, decamp. 258.6°

parahydronate m. 148.5°. The latter is accompanied by

HgCl yields 1.1 and *A*, decamp. 258.6°

73.4% *b.* *loline* decamp. 258.6°, [α]_D²⁰ 0.13°

loline m. 148.5°, decamp. 258.6°, [α]_D²⁰ 0.13°

The free base of 1 and its salt, decamp. 258.6°, [α]_D²⁰ 0.13°

b. 145.5°, m. 148.5°, decamp. 258.6°, [α]_D²⁰ 0.13°

loline m. 148.5°, decamp. 258.6°, [α]_D²⁰ 0.13°

m. 173.5°, *loline* m. 173.5°, decamp. 258.6°, [α]_D²⁰ 0.13°

The free base of 1 with *Mel* m. 219.12°

YUNUSOV, S.Yu., akademik; YULDASHEV, P.; PLEKHANOVA, N.V.

Study on alkaloids from the aboveground portion of *Vinca erecta*
Rgt. et Schmalh. Dokl. Akad. Nauk SSSR no. 7:13-15 '56.
(MIRA 12:6)

1. Akademiya nauk UzSSR (for Yunusov).
(Alkaloids) (Vinca)

YUNUSOV, S. Yu.

In the article, "Reserpinin From *Vinca erecta*," S. Yu. Yunusov, Academician of the Academy of Sciences Uzbek SSR and Director of the Institute of Chemistry, Academy of Sciences Uzbek SSR, describe the method of isolation of the alkaloid reserpinin from *Vinca erecta*, a plant of the Apocynaceae family closely related to the plant *Rauwolfia*. A total of 2.6 percent of alkaloids are extracted with ether from the roots of the plant. These include the alkaloids vinkanin-- $C_{19}H_{22}ON_2$, vinkadinin-- $C_{20}H_{24}O_3N_2$, and reserpinin-- $C_{23}H_{26}O_4N_2$; reserpinin is saponified with an alkali to form reserpinic acid. The acid and its nitrate are then methylated with diaisomethane to obtain the pure alkaloid. (Doklady Akademii Nauk Uzbekskoy SSR, No 9, 1956, pp 23-25)

Sum. 1305

YUNUSOV, S.Yu., akademik; ISMAILOV, Z.P.

Alkaloids from Linaria popovii Kuprian. Dokl.AN Uz.SSR no.11:25-27
'56. (MIRA 13:6)

1. Institut khimii AN UzSSR. 2. Akademiya nauk UzSSR (for Yunusov).
(Alkaloids) (Figwcrf)

Yunusov, S. Yu.

~~ABUDAKIROV, N.K.; YUNUSOV, S.YU.~~

Delphinium alkaloids. Zhur. ob. khim. 26 no.6:1798-1808 Je '56.
(MIRA 11:1)

1. Institut Khimii AN Uzbeckoy SSR.
(Delphonine)

YUNUSOV, S. V. - akademik, otvetstvennyy redaktor;
LYUBECHAEVSKAYA, N.I., redaktor izdatel'stva; SHEPEL'KOV, A.T.,
tekhnicheskiy redaktor

[An unknown work by al-Razi, "The Book of the Secret of Secrets."]
Neizvestnoe sochinenie ar-Razi "Kniga tainy tain." Tashkent, Izd-vo
Akad.nauk Uzbekskoi SSR, 1957. 190 p. (MIRA 10:11)

1. Akademiya nauk UzSSR (for Yunusov)
(Muhammad Ibn Zakariya, Abu Bakr, al-Razi, 10th cent.)
(Alchemy)

PAKUDINA, Z.P.; YUNUSOV, S.Yu., akademik

Thermopsis alterniflora alkaloids. Izv. AN Uz. SSR Ser. Khim,
nauk no.2:69-75 '57. (MIRA 11:8)

1. AN Uz.SSR (for. Yunusov)
(Alkaloids)

~~YUNUSOV, S.Yu., akademik; PLESKANOVA, N.V.~~

~~Study of Trichodesma incanum alkaloids. Dokl.AN Uz, SSR no.
4:31-33 '57. (MIRA 11:5)~~

1. Institut khimii rastitel'nykh veshchestv AN UzSSR.
2. AN USSR.
(Alkaloids)

YUNUSOV, S.Yu., akademik; PLEKHANOVA, N.V.

Incanine structure. Dokl. AN Uz. SSR no.5:13-16 '57. (MIRA 11:5)

1. Institut khimii rastitel'nykh veshchestv i khlopka AN UzSSR.
2. AN UzSSR (for Yunusov).

(Incanine)

YUNUSOV, S.Yu., akademik; FLEXANOVA, N.V.

Structure of trichodesmin, Dokl. AN Uz. SSR no.6:19-22 '57.

(MIRA 11:5)

1. Institut khimii rastitel'nykh veshchestv i khlopya AN UzSSR,
2. AN UzSSR (for Yunusov)

(Alkaloids)

YUNUSOV, S.Yu., akademik; AKRAMOV, S.T.; SIDYAKIN, G.P.

Study of alkaloids extracted from pabularia and hypecoum trilobium.
Dokl. AN Uz. SSR no. 7:23-25 '57. (MIRA 11:5)

1. Institut khimii rastitel'nogo syr'ya i khlopka AH UzSSR.
2. AN UzSSR (for Yunusov).

(Alkaloids)

USSR / Cultivated Plants. Medicinal. Essential Oil- M-7
Bearing. Toxins.

Abs Jour : Ref Zhur - Biologiya, No 2, 1959, No. 6478

Author : Yunusov, S. Yu.; Plekhanova, N. V.

Inst : Academy of Science, UzSSR

Title : Study of Alkaloids in Sophora Griffithii
Stock

Orig Pub : Dokl. AN UzSSR, 1957, No 8, 17-19

Abstract : Alkaloids derived from the leaves and seeds of *S. griffithii* were isolated. The content of alkaloids in leaves (~4.65%) was approximately the same during the fruit bearing stage and after the fall of fruits; 1.6% of pachycarpine and 0.57% of cytisine were obtained by separating the alkaloids. 5.93% of the alkaloids were isolated from

Card 1/2

159

~~USSR~~ / Cultivated Plants. Medicinal. Essential Oil-Bearing. Toxins. M-7

Abs Jour : Ref Zhur - Biologiya, No 2, 1959, No. 6478

seeds. 4.12% of cytisine were obtained by separating these alkaloids. The method of extraction of the alkaloids, their separation and identification are described. -- A. A. Zaytseva

Card 2/2

YUNUSOV, S.Yu.; Yuldashev, P.Kh.

Study of the alkaloids extracted from Vinca erecta Rgl. et Schmalh.
Zhur. ob. khim. 27 no. 7:2015-2018 Jl. '57. (MIRA 10:10)

1. Institut khimii rastitel'nykh veshchestv i khlopka AN Uzbekskoy
SSR.

(Alkaloids) (Apocynaceae)

AUTHORS:

Yunusov, S. Yu., Abduazimov, Kh. A.

79-12-36/43

TITLE:

An Investigation of the Four Types of Alkaloids From Ungernia
(Issledovaniye alkaloidov chetyrekh vidov Ungernia).

PERIODICAL:

Zhurnal Obshchey Khimii 1957, Vol. 27, Nr 12, pp. 3357-3361
(USSR)

ABSTRACT:

The alkaloids of the plant species Ungernia were subject to few chemical and pharmacological investigations. Only the tazettine was separated from the species "Ungernia Severtzovii" and the "likorine" from "Ungernia tadshicorum". On the investigation of the first species a production rate of 0,7 - 0,29 % of alkaloid from the bulbs was established. It was succeeded, to isolate three crystalline radicals from the alkaloid mixture of the bulbs. One of these forms a series of crystalline salts. A free alkaloid was separated from the purified chlorine hydrate. An empiric formula $C_{19}H_{23}NO_5$ of this alkaloid was computed on the basis of an elementary analysis of the radical itself as well as of its nitrate. It was called ungerine. After the removal of the ungerine from the remaining alkaloid mixture the already mentioned tazettine was separated on the basis of their different solubility in acetone and alcohol. The third alkaloid from "Ungernia

Card 1/2

An Investigation of the Four Types of Alkaloids From Ungernia 79-12-36/43

"Severtzovii" appeared to be new and was called ungeridine. It has the experimental formula $C_{20}H_{25}NO_4$ and its structure was determined more exactly (see formula!). It appears, that two new alkaloids were obtained apart from tazettine and likorine from "Ungernia Severtzovii". From the bulbs of "Ungernia tadshicorum" likorine and ungeridine were isolated, from the bulbs of "Ungernia Victoris" galamantine and likorine. The bulbs of "Ungernia ferganica" contain tazettine and likorine. These alkaloids were all four examined pharmacologically by Mushkovskiy M. D. There are 8 references, 8 of which are Slavic.

ASSOCIATION: Institute of Vegetable Raw Materials and
Cotton Chemistry (Institut khimii i rastitel'nogo syr'ya i
khlopka).

SUBMITTED: August 21, 1956

AVAILABLE: Library of Congress

1. Alkaloids - Sources

Card 2/2

YESKAIROV, M.; SIDYANIK, G.P.; YUMUSOV, S.Tu., akademik.

Alkaloids of *Haplophyllum foliosum* Vved.; folicosidine. Dokl. Akad. Nauk SSSR no. 5:23-26 '58. (NIIKA 11:8)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR.
(Bns). (Alkaloids)

SIDYAKIN, G.P.; YESKAIROV, M.; YUNUSOV, S.Yu., akademik

Alkaloids of *Haplophyllum foliosum* Vved. Structure of dubinidine.
Dokl. AN Uz. SSR no.8:27-29 '58. (MIRA 11:9)

1.Institut khimii rastitel'nykh veshchestv AN UzSSR. 2.AN UzSSR
(for Yunusov).

(Dubinidine)

SIDYAKIN, G.P.; TBSKAIROV, M.; YUNUSOV, S.Yu., akademik

Alkaloids of *Haplophyllum foliosum* Vved. Structure of dubinidine.
Dokl. AN Uz.SSR no.9:17-18 '58. (MIRA 11:12)

1. AN UzSSR (for Yunusov). 2. Institut khimii rastitel'nykh
veshchestv AN UzSSR.
(Dubinidine)

YUNUSOV, S. In., akademik; SHAKIROV, T.T.; PLEKHANOVA, N.V.

Alkaloids from *Convolvulus subhirsutus* Engl. and Schmae of the
family Convolvulaceae. Dok. AN Uz.SSR no.10:17-20 '58.
(MIRA 11:12)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR, 2. Chlen-
korrespondent AN SSSR i akademik AN UzSSR (for Yunusov).
(Alkaloids) (Bindweed)

YUNUSOV, S.Yu., akademik; PLEKHANOVA, N.V.; SHAKIROV, T.

Investigation of several species of *Bremurus*. Dokl. AN Uz.SSR
no.11:25-27 '58. (MIRA 11:12)

1. Chlen-korrespondent AN SSSR,AN UzSSR (for Yunusov). 2. Institut
khimii rastitel'nykh veshchestv AN UzSSR.
(Lilies) (Alkaloids)

YUNUSOV, S. Yu., akademik; PLEKHANOVA, N.V.

Alkaloids from *Rindera cyclodonta* Bge. from the Boraginaceae family. Dokl. AN Uz.SSR no.12:27-30 '58. (MIRA 12:1)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR. 2. AN UzSSR i chlen-korrespondent AN SSSR (for Yunusov).
(Asia, Central---Borage) (Alkaloids)

TUNUSOV, S.Yu., akademik; AKRAMOV, S.T.

* Study on alkaloids from the seeds of *Lolium cuneatum* (Novskii).
Dokl. AN Uz.SSR no.3:36-39 '59. (MIRA 12:7)

1. Institut khimii rastitel'nykh veshestv AN UzSSR. 2. AN.UzSSR
(for Yumusov). (Alkaloids)

SHAKIROV, T.; SIDYAKIN, O.P.; YUNUSOV, S.Yu., akademik

Alkaloids from seeds of *Haplophyllum perforatum*. Dokl. AN Uz.SSR
no.6:28-30 '59. (MIRA 12:9)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR. 2. AN
UzSSR (for Yumazov). (Alkaloids)

ISMAILOV, Z.F.; MAYEKH, S.Kh.; YUNUSOV, S.Yu., akademik

Alkaloids from the roots of Thalictrum simplex L. Dokl. AN Uz.
SSR no.7:32-34 '59. (MIRA 12:10)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR. 2. AN
UzSSR (for Yunusov).
(Alkaloids) (Meadow rue)

SIDYAKIN, G.P.; BESSONOVA, I.A.; YUNUSOV, S.Yu.

Alkaloids of seeds of *Haplophyllum perforatum*: Perforin. Dokl.
AN Uz.SSR no.10:33-35 '59 (MIRA 13:3)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR. 2. Chlen-korrespondent AN SSSR (for Yunusov).
(Alkaloids)

YUMUSOV, S.Yu.; ABDUSAMATOV, A.; ABDUAZIMOV, Kh.A.

Studying alkaloids of plants of the genus Jurinea. Dokl.
AN Uz.SSR no.11:29-31 '59. (MIR 13:4)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR.
2. Chlen-korr. AN SSSR (for Yunusov).
(Jurinea) (Alkaloids)

AUTHORS: Yunusov, S. Yu. and Plekhanova, N. V. SOV/79-29-2-66/71

TITLE: Alkaloids of the Plant Trichodesma Incanum (Alkaloidy Trichodesma incanum)
Structure of Incanine and Trichodesmine (Stroyeniye inkanina i trikhodesmina)

PERIODICAL: Zhurnal obshchey khimii, 1959, Vol 29, Nr 2, pp 677-684 (USSR)

ABSTRACT: Men'shikov, G. P. and Rubinshteyn, M. M. (Ref 1) separated the alkaloid trichodesmine (0.075 %) from the subterranean parts of the above plant. By lengthy extraction with ether and chloroform the authors obtained from its seeds various mixtures consisting of bases (Ref 5). On separating the alkaloid mixture (from ripe seeds) four crystalline bases were separated by making use of their different solubility in ether, benzene and acetone. The first alkaloid was unknown and was given the name of "incanine" (I); the second was the N-oxide form of incanine; the third was found to be trichodesmine (II), and the fourth was the N-oxide form of trichodesmine. Both the quantitative and qualitative composition of the alkaloids in the seeds vary markedly depending on the degree of ripeness, on the place of growth, and as far as the subterranean parts are concerned, on the plant developing stage (Table). Thus the following new

Card 1/3

Alkaloids of the Plant Trichodesma Incanum. Structure of Incanine and Trichodesmine

SOV/79-29-2-66/71

alkaloids were separated from the seeds and the upper part of the plant *Trichodesma incanum* (Bge) DC: incanine ($C_{18}H_{27}O_3N$), the N-oxide of incanine, trichodesmine ($C_{18}H_{27}O_6N$), and the N-oxide of trichodesmine. Depending on the type of saponification of incanine, new geometrical acids are formed: incanine acid and isoincanine acid from the composition $C_{10}H_{16}O_4$ (V). The conversion of the former into the latter and vice versa was put into practice. The reduction of the methyl esters of both acids with $LiAlH_4$ gave trioxy compounds having the composition $C_{16}H_{22}O_3$ (III). Compounds (V) are γ -lactone of 2-oxy-3,5-dimethylhexane-2,4-dicarbonic acid (IV). Incanine (I) has the structure of the cyclic diester of retronecine and of 2-oxy-3,5-dimethylhexane-2,4-dicarbonic acid. The structure of trichodesmine acid (VII), which is a γ -lactonic acid of 2,3-dioxy-3,5-dimethylhexane-2,4-dicarbonic acid (VI), was determined. Trichodesmine is a cyclic diester of retronecine and of 2,3-dioxy-3,5-dimethylhexane-2,4-dicarbonic acid (II). There are 1 table and 10 references, 8 of which are Soviet.

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SOV/79-29-2-66/71
Alkaloids of the Plant Trichodesma Incanum. Structure of Incanine and Trichodesmine

ASSOCIATION: Institut khimii rastitel'nykh veshchestv Akademii nauk Uzbekskoy SSR (Institute for the Chemistry of Vegetable Matter of the Academy of Sciences, Uzbekakaya SSR)

SUBMITTED: January 3, 1958

Card 3/3

5(3)

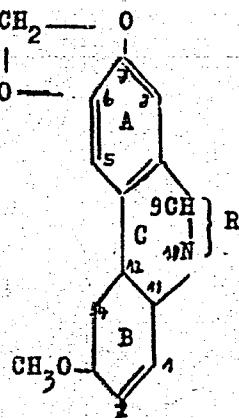
AUTHORS: Yunusov, S. Yu., Abduazimov, Kh. A. SOV/79-29-5-67/75TITLE: Investigation of the Alkaloids of *Ungernia Severtzovii*
(Issledovaniya alkaloidov *Ungernia Severtzovii*).
Structure of "Ungerine" (Stroyeniye ungerina)PERIODICAL: *Zhurnal obshchey khimii*, 1959, Vol 29, Nr 5, pp 1724-1728
(USSR)

ABSTRACT: On distilling "Ungerine" with zinc dust the authors obtained phenanthridine, and in the oxidation with potassium permanganate they obtained hydрастic acid. In the Hofmann decomposition a second double bond is formed in the tetrahydro benzene ring B, and a third by cleaving the methoxyl group as methyl alcohol; the ring is thus aromatized. This was established by the fact that benzoic acid was obtained in the oxidation of des-N-methyl "Ungerine". "Ungerine" is a derivative of phenanthridine, in which the methylene dioxy group is in position 6 -- 7. The double bond lies most probably between the carbon atoms 1 - 2, the methoxyl group in position 3.

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Investigation of the Alkaloids of *Ungernia Severtzovii*. Structure of "Ungerine"

SOV/79-29-5-67/75



There are 6 references, 2 of which are Soviet.

ASSOCIATION: Institut khimii rastitel'nykh veshchestv Akademii nauk
Uzbekskoy SSR (Institute of the Chemistry of Vegetable
Substances of the Academy of Sciences, Uzbekskaya SSR)

SUBMITTED: January 6, 1958
Card 2/2

5.3900

77414
SOV/79-30-1-75/78

AUTHORS: Sidyakin, G. P., Yeskairov, M., Yunusov, S. Yu.

TITLE: Alkaloids of the Haplophyllum Foliosum Vved.
Structure of Dubinidine

PERIODICAL: Zhurnal obshchey khimii, 1960, Vol 30, Nr 1, pp 338-
345 (USSR)

ABSTRACT:
This is a continuation of the investigation of the genus Haplophyllum A. Juss. (family Rutaceae) (Yunusov, S. Yu., Sidyakin, G. P., Zhur. Obshchey Khim., 22, 1055 (1952); 25, 2009 (1955); Doklady Akad. Nauk UzSSR, 12, 15 (1950)). The authors studied the alkaloids of the species Haplophyllum foliosum Vved. (from its stem, leaves, and green seeds). Four alkaloids were isolated: dubinidine (which was first obtained from Haplophyllum dubium Eng. Kor. -- see the reference cited above), skimmianine, and two new alkaloids called by the authors "foliosine" (foliozin) (Doklady Akad. Nauk

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Alkaloids of the Haplophyllum Foliosum Vved.
Structure of Dubinidine

77414

SOV/79-30-1-75/78

UzSSR, 2, 21 (1957) and "foliosidine" (foliozidin) (ibid., 5, 23 (1958)). Dubinidine was precipitated from aqueous solution of the alkaloids (this solution was obtained by treating the chloroform extracts of *H. foliosum* with sulfuric acid) with gaseous NH₃.

The precipitate was dissolved in acetone and converted into the hydrochloride (mp 195-196°, [α]_D¹⁸ (-53.92°)).

Pure dubinidine (mp 132-133°, [α]_D^{26.5} (-62.95°)) was obtained by addition of concentrated NH₄.OH to aqueous suspension of its hydrochloride. Its ultra-violet spectrum is shown in the figure below, along with spectrum of dihydroflindersine (which has a pyranoquinoline structure).

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Alkaloids of the Haplophyllum Foliolum
Vved. Structure of Dubinidine

77414
SOV/79-30-1-75/78

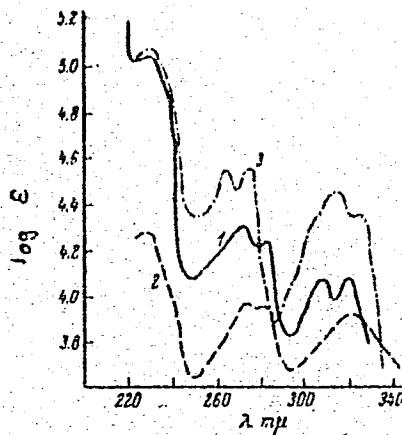


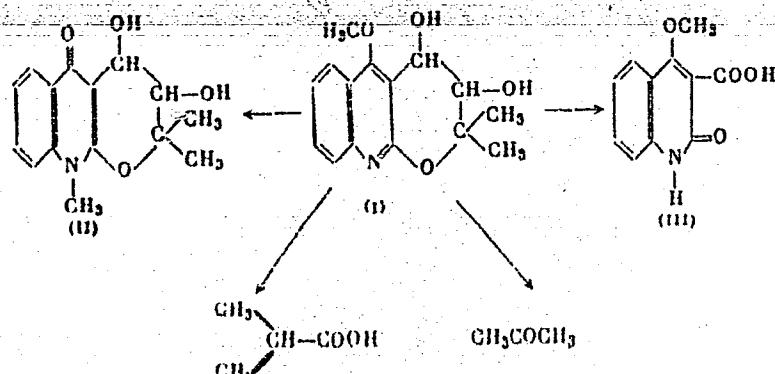
Figure. Ultraviolet absorption spectra. (1)
dubinidine; (2) dihydroflindersine; (3) 4-methoxy-
quinoline-2.

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Alkaloids of the Haplophyllum Foliosum
Vved. Structure of Dubinidine

77414
SOV/79-30-1-75/78

Decarboxylation, iodomethylation, oxidation (with KMnO₄, chromic, and periodic acids) etc., have proven the structure of dubinidine to be identical with (I) in the figure below, i.e., with 2,2-dimethyl-3,4-dioxy-5-methoxy- α , β -dihydropyranoquinoline.



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Alkaloids of the Haplophyllum Foliosum
Vved. Structure of Dubinidine

77414
SOV/79-30-1-75/78

The following derivatives of dubinidine were prepared and investigated: hydrobromide (mp 197-198°); hydroiodide (mp 161-162°, $[\alpha]_D^{18}$ (-47.32°)); nitrate (mp 176-177°, $[\alpha]_D^{22}$ (-52.39°)); methiodide mp 153-154° which, upon addition of alcoholic alkali, gave isodubinidine, compound (II) in the figure above (mp 214-215°, $[\alpha]_D^{25}$ (+21.05°)); diacetyldubinidine (mp 108-109°, $[\alpha]_D^{19}$ (-47.70°)). Oxidation with $KMnO_4$ led to an aldehyde and then to the optically inactive dictamninic acid (III in the figure above). Skimmianine was separated from the other two alkaloids (the solid mixture was obtained from the chloroform extracts of the solution which was left after precipitation of dubinidine by triturating the mixture in acetone, which dissolves foliosine and foliosidine). The mixture of the two latter compounds was purified by subsequent addition of 10%

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Alkaloids of the Haplophyllum Foliosum
Vved. Structure of Dubinidine

77414
SOV/79-30-1-75/78

HCl and ammonia and extraction with chloroform (followed by distillation of the latter). The residue was dissolved in methanol and acidified with alcoholic HCl. The optically inactive foliosine hydrochloride (mp 253-254°) fell out after addition of threefold amount of acetone to the cooled solution and was converted to foliosine by addition of 25% NH₄OH (mp 188-189°). The following foliosine derivatives were prepared: hydrobromide (mp 249-250° (decomp.)); hydroiodide (mp 225-226° (decomp.)); nitrate (mp 170-171.5° (decomp.)); methiodide (mp 210-211°); and perchlorate (mp 229-231° (decomp.)). Its formula was found to be: C₁₅H₁₀O(NCH₃)₂(CH₂O₂).

The alkaloid residue isolated from the acetone solution, which was left after precipitation of foliosine, contained mainly folisidine, C₁₇H₂₃O₅N, mp 141-142° and [α]_D²⁵ (+41.62°). Its more detailed formula was found to be C₁₅H₁₅O₂(NCH₃)(OCH₃)(OH)₂.

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Alkaloids of the Haplophyllum Foliosum
Vved. Structure of Dubinidine

77414
SOV/79-30-1-75/78

The ultraviolet spectrum (in alcohol) has the following maxima: $\lambda_{\text{max}}^{234}$ ($\log \epsilon$, 4.94); 252 ($\log \epsilon$, 4.92); 324 ($\log \epsilon$, 3.98); 234 m μ ($\log \epsilon$, 3.84). Abstracter's Note: Two maxima $\lambda_{\text{max}}^{234}$ are

given in the article. The following derivatives: were prepared: hydrobromide (mp 167-168°); hydrochloride (mp 162-164°); foliosidine picrate (mp 182-183°); and diacetylfoliosidine (mp 129-130°, $[\alpha]_D^{18}$ (+14.95°)). There is 1 figure; and 14 references,

erences, 7 Soviet, 5 German, 1 U.K., and 1 U.S. The U.S. and U.K. references are: G. Sidney, A. F. Smith, E. C. Horning, J. Am. Chem. Soc. 79, 2239 (1957); R. F. C. Brown, J. J. Hoobs, L. K. Huges, E. Ritchie, Austral. J. Chem., 7, 4, 348 (1954).

ASSOCIATION: Institute of Chemistry of Plant Substances, Academy of Sciences of the UzbekSSR (Institut khimii rastitel' Card 7/8

Alkaloids of the Haplophyllum Foliosum
Vved. Structure of Dubinidine

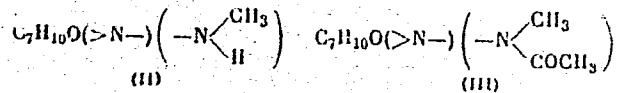
77414
SOV/79-30-1-75/78

nykh veshchestv Akademii nauk Uzbekskoy SSR)

SUBMITTED: October 29, 1958

Card 8/8

5.3610, 5.3900

77918
SOV/79-30-2-69/76**AUTHORS:** Yunusov, S. Yu. Akramov, S. T.**TITLE:** Investigation of Alkaloids of Lolium Cuneatum.
Communication II.**PERIODICAL:** Zhurnal obshchey khimii, 1960, Vol 30, Nr 2,
pp 677-682 (USSR)**ABSTRACT:** The authors reported previously (this j., 1955,
Vol 25, p 1813) the separation of 3 new alkaloids
from the seeds of *Lolium cuneatum* Neyski (Fam.
Gramineae), which they named lolimidin (I), lolin
(II), and lolinin (III).

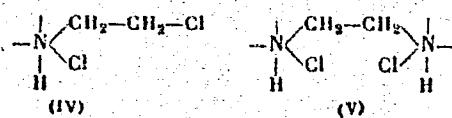
Card 1/5

It was also established (Izv. AN UzSSR, ser. khim.,
1957, Vol 2, p 69) that dichloroethane condenses
readily with alkaloids containing primary and

Investigation of Alkaloids of *Lolium Cuneatum*. Communication II.

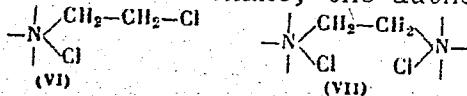
77918
SOV/79-30-2-69/78

secondary nitrogen atoms in the molecule, and that it forms type (IV) or (V) compounds. Dichloroethane can form also type (VI) or (VII) compounds on condensation with strong tertiary bases.



The yield of IV-VII depends on the time and temperature of the reaction. Lolin II was condensed with chloroethane and gave crystalline compounds of IV or VI structure (yield 65%; mp 135-136°C) named by the authors lolinchloroethane. Taking into account the above properties of dichloroethane, the authors extracted

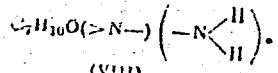
Card 2/5



Investigation of Alkaloids of Lolium
cuneatum. Communication II.

SGV/79-30-P-LJ/76

Lolium cuneatum seeds with chloroform; the extract, after separation of substances soluble and insoluble in acetone, gave, on treatment with a methanolic solution of HCl and a methanolic solution of sodium perchlorate, a new alkaloid named norlololin (VIII), obtained in the form of its diperchlorate. Free norlololine had a bp 94-95° C at 2 mm; d_{20}^{20} 1.1793; n_p^{16} 1.5220; on standing, it absorbed CO_2 from air and gave a crystalline carbonate (mp 141° C).



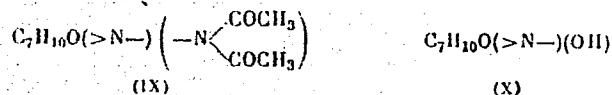
It was shown that VIII is identical with the product of oxidation of lolin with KMnO_4 in an acid medium, and that it gives easily dinitrates, dihydrochlorides and dipicrates. Acetylation of VIII gave N-diacetyl-norlololin (IX) bp 190-195° C at 2 mm, a glass-like, noncrystallizable substance, which on heating with

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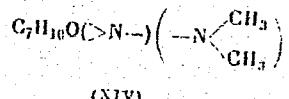
Investigation of Alkaloids of *Lolium Cuneatum*. Communication II.

77918
SOV/79-30-2-69/7

30% sulfuric acid gave again VIII. Diazotization of VIII gave a crystalline amino alcohol (X; mp 192° C), named by the authors heminorlolin, which gave readily the corresponding hydrochloride, bromohydrate, and picrate.



Methylation of VIII with formaldehyde and formic acid in 1:2:2 molar ratio gave N-methylloolin (XIV), which with KMnO_4 and sulfuric acid (2 g-atoms O), was oxidized to II; similar oxidation in acetone (6 g-atoms O) gave VIII.



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Investigation of Alkaloids of *Lolium cuneatum*. Communication II.

77918

SOV/79-30-2-69/78

Lolin II was oxidized to VIII with KMnO₄ in sulfuric acid. There are 7 Soviet references.

ASSOCIATION: Chemical Institute for Plant Substances, Academy of Sciences of the Uzbek SSR (Institut khimii rastitel'nykh veshchestv Akademii nauk Uzbekskoy SSR)

SUBMITTED: February 25, 1959

Card 5/5

5.3610,5.3900

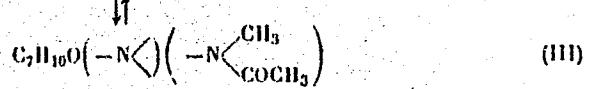
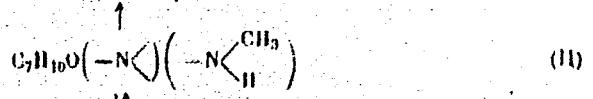
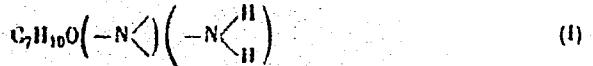
77919
SOV/79-30-2-70/78

AUTHORS: Yunusov, S. Yu., Akramov, S. T.

TITLE: Investigation of Alkaloids of Lolium Cuneatum. Communication III

PERIODICAL: Zhurnal obshchey khimii, 1960, Vol 30, Nr 2, pp 683-689 (USSR)

ABSTRACT: Norlololin (I), lolin (II), and lolinin (III) have a common heterocyclic ring; this has been demonstrated in the preceding study by the mutual conversion of one of the above alkaloids into another (our abstract 77918).

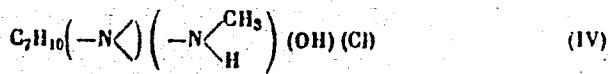


Card 1/3

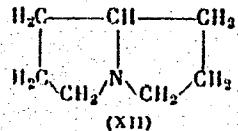
Investigation of Alkaloids of Lolium
Cuneatum. Communication III

77919
SOV/79-30-2-70/78

Lolin on heating with 15% HCl in sealed ampoules at
130-140° C gave hydroxychlorololin (IV; mp 105-106° C
from acetone; $[\alpha]_D^{16} -74.49$).



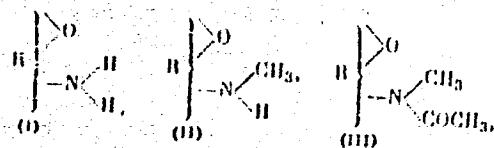
It was established that the properties of I-III coincide
closely with the properties of pyrrolizidine (XII):



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Investigation of Alkaloids of *Lolium Cuneatum*. Communication III77919
SOV/79-30-2-79/78

Hence, the above 3 alkaloids are derivatives of XII and can be represented by the formulas:



where R is the radical obtained by subtracting 3 hydrogen atoms from the XII molecule. There are 5 references, 1 U.S., 4 Soviet. The U.S. reference is: N. G. Brink, F. A. Kuehl, Jr., E. H. Flunn, J. Am. Chem. Soc., 68, 2557 (1946).

ASSOCIATION: Chemical Institute for Plant Substances, Academy of Sciences of Uzbek SSR (Institut khimii rastitel'nykh veshchestv Akademii nauk Uzbekskoy SSR)

SUBMITTED: March 16, 1959

Card 3/3

YUNUSOV, S.Yu.; ISMAILOV, Z.F.

Alkaloids of Thalictrum minus L, Part 3: Structure of thalmine. Zhur. ob. khim. 30 no.5:1721-1727 May '60.
(MIRA 13:5)

1. Institut khimi rastitel'nykh veshchestv Akademii nauk
Uzbekskoy SSR.
(Alkaloids)

YUNUSOV, S.Yu.; AKRAMOV, S.T.

Structure of norloline, loline, and lolinine. Part 4. Zhur. ob.
khim. 39 no.9:3132-3137 S '60. (MIRA 13:9)

Il Institut khimii rastitel'nykh veshchestv Akademii nauk Uzbekskoy
SSR.
(Loline) (Lolinine) (Norloline)

AEDUSAMATOV, A.; AEDUAZIMOV, Kh.A.; YUNUSOV, S.Yu.

Alkaloids from *Ungernia victoris* VVED. Uzb.khim.zhur. 6
no.1:45-55 '62. (MIRA 15:3)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR.
(Alkaloids)

SIDYAKIN, G. P.; BESSONOVA, I. A.; PASTUKHOVA, V. I.; XUNUSOV, S. Yu.

Alkaloids Haplophyllum. Part 3: Structure of dubinidine and
dubamine. Zmtr. ob. khim. 32 no.12:4091-4096 D '62.
(MIRA 16:1)

1. Institut khimii rastitel'nykh veshchestv AN Uzbekskoy SSR.

(Alkaloids) (Dubinidine)

YULDASHEV, P.A.; YUNUSOV, S.Yu.

Structure of vincanine. Uzb.khim.zhur. 7 no.1:44-49 '63.
(MIRA 16:4)

1. Institut rastitel'nykh veshchestv AN UzSSR.
(Vincanine)

UBAYEV, Kh.; YULDASHEV, P.Kh.; YUNUSOV, S.Yu.

Study of alkaloids of Pedicularis olgae RGL. Uzb.khim.zhur. 7 no.3:
33-36 '63. (MIRA 16:9)

Institut khimii rastitel'nykh veshchestv AN UzSSR.
(Figwort) (Alkaloids)

FAXHRUTDINOVA, I.M.; SIDYAKIN, G.P.; YUNUSOV, S.Yu.

Alkaloids from Haplophyllum folosium. Haplopholin. Uzb. khim.
zhur. 7 no.4:41-43 '63. (MIRA 16:10)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR.

ABDUSAMATOV, A.; ABUDAZIMOV, Kh.A.; YUNUSOV, S.Yu., akademik

Alkaloids from Ungernia tadshicorum Vved. and artificial alkaloids
from Ungernia victoris Vved. Dokl. AN Uz. SSR 20 no.1:18-21 '63.
(MIRA 16:6)

1. Institut khimii rastitel'nykh veshchestv AN Uzbekskoy SSR.
2. AN Uzbekskoy SSR (for Yunusov).
(Alkaloids) (Ungernia)

ABDUAZIMOV, Kh.A.; YUNUSOV, S.Yu.

Structure of ungerine. Dokl. AN SSSR 153 no.6:1315-1317
D '63. (MIRA 17:1)

1. Institut khimii rastitel'nykh veshchestv AN U2SSR.
2. Chlen-korrespondent AN SSSR (for Yunusov).

ALLAYAROV, Kh.; ABDUAZIMOV, Kh.A.; YUNUSOV, S.Yu.

Alkaloids of Ungernia triphaera EGE. Uzb.khim.zhur. 8 no.2:
46-51 '64. (MIRA 17:5)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR.

HESSONOVА, I.A.; SIDYAKIN, G.P.; YUNUSOV, S.Yu.

Alkaloids of Haplophyllum dubium. Structure of dubinine. Zhur.ob.
khim. 34 no.1:347-351 Ja '64. (MIRA 17:3)

1. Institut Khimii rastitel'nykh veshchestv AN UzSSR.

SMIRNOVA, L.S.; ABDUAZIMOV, Kh.A.; YUNUSOV, S.Yu.

Alkaloids of *Ungernia severtzovii*. Structure of unsevine. Dokl.
AN SSSR 154 no.1:171-173 Ja'64. (MIRA 17:2)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR. Chlen-
korrespondent AN SSSR (for Yunusov).

YULDASHEV, P.Kh.; YUNUSOV, S.Yu.

Vincarine, a new alkaloid from the roots of Vinca erecta RGL. et Schmalh.
(MIRA 17:2)
Dokl. AN SSSR 154 no.6:1412-1413 F '64.

1. Institut khimi rastitel'nykh veshchestv AN UzSSR. 2. Chlen-korrespondent AN SSSR (for Yunusov).

MAYEKH, S.Kh.; YUNUSOV, S.Yu., akademik

Alkaloids of Thalictrum simplex L.; structure of talsimin.
Dokl. AN Uz.SSR 21 no.9:27-29 '64. (MIRA 19:1)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR.
2. Akademiya nauk UzSSR (for Yunusov).

SMIRNOVA, L.S.; ABDUAZIMOV, Kh.A.; YUNUSOV, S.Yu.

Alkaloids of *Ungernia sovertzowii* (Rga.) B. Fed. Khim. prirod.
sozdr. no. 5: 322-328. '65. (MIRA, 18:12)

1. Institut khimii rastitel'nykh veshchestv AN UeSSR.
Submitted April 5, 1965.

LUTFULLIN, K.L.; YULDASHEV, P.Kh.; YURUSOV, S.Yu.

Study of the alkaloids of Pedicularis oligae. Structure of plantagonin and indicain. Khim. prirod. soed. no.5:365-366 '65.
(MIRA 18:12)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR. Submitted
August 6, 1965.

KUCHENKOVA, M.A.; YULDASHEV, P.Kh.; YUNUSOV, S.Yu.

Vinervine, a new alkaloid from the above-ground part of
Vinca erecta RGL et Schmalh. Izv. AN SSSR. Ser. khim. no. 12:2152-
2155 '65. (MIRA 18:12)

1. Institut khimii ras'sitel'nykh veshchestv AN UzSSR.
Submitted July 29, 1963.

SHAKIROV, R.; NURIDDINOV, R.N.; YUNUSOV, S.Yu.

Synthesis of "edpetilin." Dokl. AN SSSR 161 no.3:620-621 Mr '65.
(MIRA 18:4)

1. Institut khimii rastitel'nykh veshchestv AN Uzbekskoy SSR.
2. Chlen-korrespondent AN SSSR (for Yunusov).

TELEZHENETSKAYA, M.V.; YUNUSOV, S.Yu.

Structure of thalmine and thalmidine. Dokl. AN SSSR 162 no.2;
354-355 My '65. (MIRA 18:5)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR. 2. Chlen-korrespondent AN SSSR (for Yunusov).

YUNUSOV, S.Yu.; MNATSAKANYAN, V.A.; AKRAMOV, S.T.

Alkaloids of some species of Papaver and Roemeria and the structure
of fugapavin. Izv. AN SSSR. Ser. khim. no.3:502-509 '65. (MIRA 18:5)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR,

YUNUSOV, M.S.; AKRAMOV, S.T.; YUNUSOV, S.Yu.

Alkaloids of Corydalis gortschakovi and Corydalis pseudocaudunga.
Dokl. AN SSSR 162 no. 3:607-609 My '65. (MIRA 18:5)

1. Institut khimii rastitel'nykh veshchestv AN UzSSR. 2. Chlen-korrespondent AN SSSR (for S.Yu.Yunusov).

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